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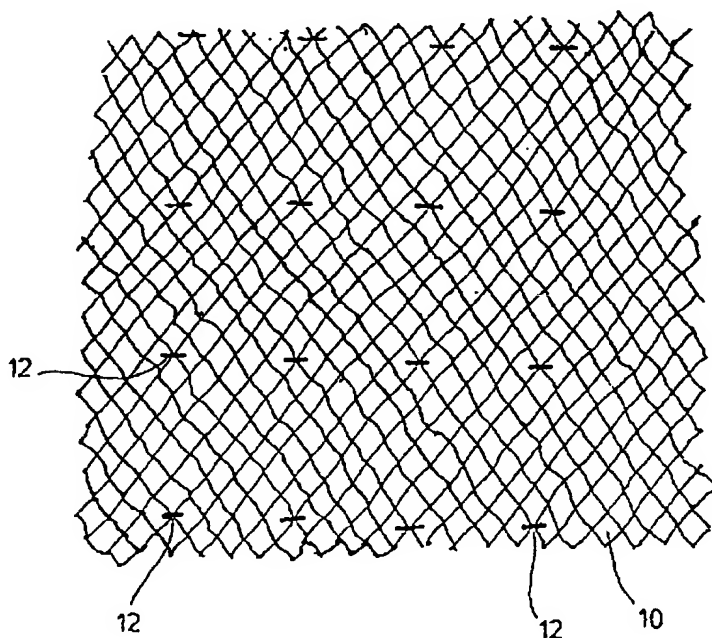
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(54) Title: **AREAL IMPLANT WITH X-RAY-VISIBLE ELEMENTS**



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(57) Abstract: An areal implant has a flexible polymer-based basic structure (10) and X-ray-visible elements (12) and is preferably set up as a permanent implant. The X-ray-visible elements can be arranged in an areal pattern in order to be able to judge the position of the implant in the patient at any time by means of an X-ray procedure.

Areal implant with X-ray-visible elements

The invention relates to an areal implant with a flexible polymer-based basic structure.

Areal implants with a flexible polymer-based basic structure, which are prepared for example in the form of meshes or tapes, are widely used. They are used for example in a surgical operation, in order to support or to reinforce an organ or tissue or to promote the healing procedure. Such an implant often has to remain permanently or at least for a very long time in the body of a patient. In this case, the basic structure contains non-resorbable polymer or only very slowly-resorbable polymer.

In the course of time, the inserted implant can shift, shrink or fold. This can lead to problems for the patient. Diagnostically, by means of imaging procedures, this can be detected, if at all, only with great difficulty, as conventional areal implants are relatively fine, in order to ensure a sufficient flexibility, and have already become fused with tissue a short time after the operation, so that they can hardly be made out, if at all, using conventional and widespread diagnostic methods such as ultrasound or X-ray procedures, so that no diagnostically usable conclusions are possible.

Metal meshes can be made visible with X-ray procedures, but are not sufficiently flexible.

In EP 0 894 481 temporary and removable X-ray-visible markers are described, which are essentially applied on stents. A stent is generally taken to be a tubular metal mesh or a perforated metal tube which is much more inelastic than an areal implant with a flexible basic structure on polymer base. The markers shown in this document are not suitable for an areal implant which is permanently inserted in the body of a patient.

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A temporary marking with a resorbable marker which, as a rule, consists of a resorbable polymer and an element such as zirconium or iodine, is described in EP 0 894 503. The use of a resorbable and physiologically compatible polymer which is combined
5 with toxic elements such as elementary zirconium or iodine, is problematical.

In WO 94/01056 stent implants are shown which consist of a tubular metal mesh and loop-shaped metal wires knotted onto them
10 which are made from an X-ray-dense metal, such as tantalum. These implants are likewise relatively bulky, as they are meant to support vessels, and are not comparable with areal implants with a flexible polymer-based basic structure.

15 R. Sahagian (Critical Insight: Marking Devices with Radioopaque Coatings, MD&DI May 1999) describes X-ray-visible coatings on stents and catheters which are produced with the help of a sputtering process. These coatings are intended for short-term implants such as catheters or for not very flexible metal meshes
20 such as stents and are unsuitable for permanently implanted flat polymer implants. Furthermore, sputtering processes have the disadvantage that often only a fraction of the expensive precious metals used is deposited on the implant.

25 Swabs marketed by the firm Hartmann under the name "Telacomp", contain a polypropylene thread filled with barium sulphate. However, these swabs are not proposed for permanent implantation. The marking only reveals whether a swab has remained in the body after an operation, but gives no information about the
30 orientation of the swab in the body, as the thread filled with barium sulphate runs only in the middle of the swab and thus a shifting or a folding at the edge cannot be recognised. Barium sulphate is to be considered as critical for use in a long-term implant, because of the toxicity of barium ions, if it is not
35 sufficiently encapsulated.

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The object of the invention is to create a possibility, which is economical and safe for the patient, of being able to monitor a flexible areal implant which is implanted in the body of a patient for a long time or permanently, after the operation at any
5 chosen time, in a way which does not burden the patient.

This object is achieved by an areal implant having the features of claim 1. Advantageous designs of the invention are given in the dependent claims.

10

The areal implant according to the invention has a flexible polymer-based basic structure and has X-ray-visible elements. The implant is preferably set up for permanent implantation, the X-ray-visible elements being present in tissue-compatible form,
15 i.e. if at all possible releasing no toxic substances even after a long time, and being permanently connected to the basic structure. The implant is preferably flexible as a whole. The X-ray-visible elements enable the implant to be made visible as required at any time after the surgical operation to insert the im-
20 plant. It is particularly advantageous to have the X-ray-visible elements arranged in an areal pattern. This is because, in this case, a shift of the implant or of sections of the implant (e.g. folding of corner) can be easily recognised on the X-ray image. Shrinking or stretching is also visible through the changed
25 distances between the individual components of the pattern.

The basic structure preferably has a non-resorbable polymer so that it is suitable for a permanent implant, but can also have a proportion of resorbable polymer. Examples of tissue-compati-
30 ble non-resorbable or very slowly resorbable substances are polyalkenes (e.g. polypropylene or polyethylene), fluorinated polyolefins (e.g. polytetrafluoroethylene or polyvinylidene fluoride), polyamides, polyurethanes, polyisoprenes, polystyrenes, polysilicones, polycarbonates, polyaryletherketones (PEEK),
35 polymethacrylates, polyacrylates, aromatic polyesters, polyimides as well as mixtures and/or copolymers of these substances.

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Polyhydroxy acids (e.g. polylactides, polyglycolides, polyhydroxybutyrates, polyhydroxyvalerates), polycaprolactones, polydioxanones, synthetic and natural oligo- and polyaminoacids, polyphosphazenes, polyanhydrides, polyorthoesters, polyphosphates, 5 polyphosphonates, polyalcohols, polysaccharides, polyethers, resorbable glasses as well as mixtures and/or copolymers of such substances, for example, come into consideration as resorbable substances.

10 The flexible basic structure is preferably designed as mesh, tape, foil or perforated foil and in principle it can be of conventional type. It is preferably thinner than 1 mm. It is conceivable that the shape of the implant to be used in a given surgical operation is cut to size from a larger piece of material 15 before the operation.

X-ray-visible elements, which are particularly clearly visible in an X-ray procedure, contain a chemical element or several chemical elements of a medium or high atomic number in sufficient density. 20 Non-toxic and chemically stable chemical elements or chemical compounds with these properties are particularly suitable. If no sufficient long-term stability is guaranteed, such as for example in the case of barium sulphate, additional measures are recommended, as explained further below. Examples 25 of X-ray-visible substances which can be used in the X-ray-visible elements are pure zirconium dioxide, stabilized zirconium dioxide, zirconium nitride, zirconium carbide, tantalum, tantalum pentoxide, barium sulphate, silver, silver iodide, gold, platinum, palladium, iridium, copper, ferric oxides, not very 30 magnetic implant steels, non-magnetic implant steels, titanium, alkali iodides, iodated aromatics, iodated aliphatics, iodated oligomers, iodated polymers as well as mixtures and alloys of such substances. Non-magnetic materials offer advantages because, for example, they do not disturb imaging diagnostic procedures 35 using magnetic resonance.

- 5 -

In a preferred version of the invention, at least part of the X-ray-visible elements is formed as pre-shaped bodies of respective length, width and height in the range 0.1 mm to 50 mm, the pre-shaped bodies being attached to the basic structure.

5

The pre-shaped bodies can be present in many different shapes, e.g. as beads, balls, small tubes, rods, small plates, rings, discs, bones or clips. A coating or outer wrapper (preferably made from polypropylene or beeswax) made from a non-resorbable material can be advantageous, in particular if the long-term stability of an uncoated pre-shaped body is doubtful. Such a pre-shaped body can be permanently connected to the flexible basic structure in many different ways, in particular by knotting, compression, welding and/or bonding (e.g. with or with the help of foils, small foil strips, foil tubes, small meshes or adhesives), by attachment direct to the basic structure or to a holding device (such as e.g. a thread, a mesh, etc) connected to the basic structure.

20 The pre-shaped bodies (scattering bodies) can be designed and connected to the basic structure in many different ways. Citable as further examples are perforated bodies (e.g. sleeves, rings, beads) which are strung on a thread or attached to a thread and consist of an X-ray-visible substance, for example zirconium dioxide, barium sulphate, non-magnetic implant steel, titanium, gold or other precious metals. In addition, the pre-shaped bodies can be coated with one of the polymers used for the basic structure. In a preferred version, at least one pre-shaped body contains a mixture of at least one X-ray-visible substance with a binding agent, preferably a non- or slowly resorbable polymer and/or a wax. Furthermore, the X-ray-visible substances can be encapsulated in a glass. Pre-shaped bodies attached to a thread can be directly incorporated into the flexible basic structure during the manufacture of the implant or fixed on the basic structure by suitable thread connections or thermal fixing.

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The pre-shaped bodies considered have the advantage that, because of their size, they can be also made visible with other diagnostic procedures such as ultrasound methods and magnetic resonance procedures. Furthermore, the flexibility of the implant is not, or is only slightly, impaired by the presence of the pre-shaped bodies. A correspondingly thick X-ray-visible monofilament thread or wire, the X-ray visibility of which corresponds to that of the pre-shaped bodies present at the implant, would on the other hand lead to a marked stiffening of the flexible basic structure, which is not desired.

Pre-shaped bodies designed as clips are particularly suitable for subsequent attachment to a flexible basic structure. Metal clips are considered for example, but also clips which are cast from a composite material which contains X-ray-visible substances in the form of chemical elements, oxides, salts or organic compounds which are compounded in a non-resorbable polymer.

A pre-shaped body can also be connected to the basic structure in an advantageous manner by covering it with thin foil pieces on both sides and attaching it to the basic structure via two-sided welding with ultrasound.

In a preferred design of the invention at least some of the X-ray-visible elements have a polymer tube or a cord which is at least partly filled with particles of a size of at most 2.5 mm made from an X-ray-visible substance. For preference, the X-ray-visible substance is additionally fixed in the polymer tube or the cord, for example by thermal shrinking and/or gluing.

With the help of such a polymer tube or such a cord, an X-ray-visible substance can be introduced into the basic structure as a quasi-linear element. The implant is preferably so designed that the flexibility, the elasticity, the rigidity and the tensile strength are not adversely affected.

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A particularly preferred type of attachment of the X-ray-visible elements to the basic structure is one which can be used for pre-shaped bodies in particular, and in which a pre-shaped body or several pre-shaped bodies (e.g. balls made from zirconium dioxide) are packed into short tubes, e.g. a few centimetres long, at equal intervals. If the basic structure is a mesh made from polypropylene, these tubes are preferably made from polyethylene or polypropylene for medical use and have, in the lumen, an equal or somewhat greater diameter than the pre-shaped bodies. These short, filled tubes are thermally fixed on the mesh, optionally under pressure. Short strips thus form, in which the X-ray-visible substance is completely encapsulated and which are so strongly fixed that these strips or the enclosed particles of the X-ray-visible substance cannot be removed from the mesh without destroying it. The mechanical properties of the basic structure, such as for example the flexural strength, are not adversely affected.

In a further preferred version of the invention, at least part of the X-ray-visible elements is formed as polymer, into which an X-ray-visible substance made from particles of a size in the range of 10 nm to 500 μm (preferably 10 nm to 100 μm) is compounded.

The polymer with X-ray-visible substance can be present in the implant for example in the form of monofilaments, filaments reduced along sections, multifilaments, twines, braided threads, cords, foils, films, small film tapes, tapes or such forms which are provided with knots. In advantageous designs, the basic structure has polymer with X-ray-visible substance, which for example can be incorporated direct into the basic structure (e.g. knitted in), or the polymer with X-ray-visible substance is attached to the basic structure as additional component; mixed forms are also conceivable. Preferred examples of the X-ray-visible substance of the polymer are zirconium dioxide as well as barium sulphate; in the latter case the polymer should

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be coated in addition with a non-resorbable polymer or wax, in order to prevent the barium sulphate, which has a toxic effect, from being released in the body of a patient in the long term.

5 Thus, for example, zirconium dioxide particles with an average diameter of less than 1 μm can be compounded particularly advantageously in polypropylene with a mass content of 10% to 90%, preferably 50%, and mono- and multifilaments can be extruded from this material. These can be incorporated into the basic
10 structure in the form of threads, cords or thin tapes, so that cutting is possible for the surgeon, without the X-ray-visible elements coming loose from the supporting basic structure.

In the case of monofilament threads, care is to be taken that
15 the threads, when they have good X-ray visibility, are also still sufficiently elastic in order to not adversely affect the flexural properties of the implant. This can be achieved by diluting a relatively thick X-ray-visible thread with a diameter of for example 0.2 mm to 2 mm at pre-set intervals by thermal
20 treatment and stretching and optionally reacting it with plasticizers or plasticizing polymers (e.g. adding of polyethylene to polypropylene). Such X-ray-visible filaments are preferably incorporated into the basic structure so that they have no supporting function and cannot adversely affect the properties of
25 the implant, which are essentially dictated by the basic structure, such as its tensile strength or elasticity.

Monofilaments or multifilaments made from polymer with X-ray-visible substance can be knotted once or repeatedly in the area
30 of the basic structure. The local visibility in the X-ray image is thereby increased, without the flexural strength of the filaments being increased to an extent such as that which is present in a thread with the knot thickness. Furthermore, as a result of their size, such knots can also be detected with other diagnostic
35 stic methods, such as ultrasound procedures or magnetic resonance.

It is particularly advantageous if at least part of the X-ray-visible elements has an X-ray-visible symbol which is preferably provided repeatedly and at equal intervals. Such a symbol can be sewn for example from X-ray-visible threads, stitched from X-ray-visible threads, embossed from X-ray-visible foil or put together from X-ray-visible objects or X-ray-visible powder.

The symbol is preferably neither point- nor mirror-symmetrical; thus the lower-case letter "e" can, for example, be considered.

10 With the help of the symbol or the symbols, a deformation of the implant, e.g. an undesired knotting or overlapping, on the one hand from the relative distances between the symbols and on the other hand from a change in direction of the symbols can be recognised in the X-ray image. It is also conceivable to use the

15 product name of the implant or the company name of the manufacturer as the symbol, e.g. in the form of embossed pieces made from X-ray-visible foil, which are attached at equal intervals to the basic structure.

20 It is of particular advantage if the implant in the body of the patient can also be detected with the help of ultrasound procedures and/or in magnetic resonance tomography. This can be achieved in particular with the help of larger pre-shaped bodies or knots, as already mentioned.

25 It is thus possible, using the invention, to provide implantable flexible polymer meshes and tapes with an X-ray-visible substance so that the desired properties, such as low weight, elasticity, tensile strength and behaviour at the implantation site, are

30 not essentially altered and also other diagnostic procedures, such as X-ray photographs, computer tomography photographs of regions lying behind the implant or magnetic resonance procedures, are not noticeably disturbed by the marking with X-ray-visible elements. Furthermore, it is possible to anchor the

35 marking so firmly to the basic structure so that no false diagnoses are effected by a migration of loosened X-ray-visible

- 10 -

elements. Furthermore, the implant material can be marked such that it is possible for the doctor to cut it to size later without causing a loosening of the X-ray-visible elements, whereby size, position and shape of the implant would no longer be able to be detected with certainty. The marking with the X-ray-visible elements can be designed to be toxicologically harmless, and this to be effective for decades. Thus, not only proven inert substances such as e.g. gold or titanium (which develops a protective layer) are suitable, but also, in sufficiently and permanently encapsulated form e.g. barium sulphate, which is otherwise only permitted as an oral X-ray contrast medium because barium ions are very toxic and, despite the low solubility of barium sulphate of 2.5 µg/ml, toxic effects can be expected in the case of long-term implantation (Chang, Biomaterials 2, 151-155, 1981).

The invention is illustrated in the following with the help of individual embodiments. The drawings show in

- 20 Figure 1 a top view of the implant according to Example 1, which has clips made from titanium as X-ray-visible elements,
- 25 Figure 2 a top view of the implant according to Example 9, which has balls made from zirconium dioxide as X-ray-visible elements,
- 30 Figure 3 a schematic representation of the implant according to Example 14, which is produced on a crochet galloon machine and has X-ray-visible threads, and
- Figure 4 a thread course representation to illustrate the manufacture of the implant according to Example 15.

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Example 1

Manufacture of an X-ray-visible, fine and coarse-pored polypropylene mesh with clips made from titanium

5 A partly-resorbable implant mesh customary in the trade, marketed by Ethicon GmbH under the name "Vypro", was boiled in a 10% soda solution, rinsed with water and air-dried in order to remove the resorbable part. Clips made from titanium ("LIGACLIP Extra", small, manufactured by Ethicon Endo-Surgery) were attached to the resulting fine, but coarse-pored polypropylene mesh
10 on the intersection points of every seventh and eighth wale using the associated applicator. Figure 1 shows the basic structure 10 in the form of the polypropylene mesh as well as the clips designated by 12.

15 The implant made in this way was placed under a 10 cm-thick gel cushion for sonography, in order to have an absorption comparable to the in-vivo-situation, and X-rayed (focus-film distance: 1m, exposure 52/2.5). The X-ray-visible elements in the form of
20 clips 12 were clearly visible in the X-ray image. They could not be separated from the mesh by gentle manual pulling.

Example 2

Manufacture of an X-ray-visible, fine and coarse-pored polypropylene mesh with clips made from implant steel
25

A partly-resorbable implant mesh customary in the trade ("Vypro", Ethicon GmbH) was boiled in a 10% soda solution, rinsed with water and air-dried, in order to remove the absorbable
30 part. Implantable clips made from steel ("LIGACLIP Extra", small, stainless steel with the US-material number 316L, from Ethicon Endo-Surgery) were attached to the resulting fine, but coarse-pored polypropylene mesh (basic structure) on the intersection points of every seventh and eighth wale using the associated applicator.
35

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The implant made in this way was placed under a 10 cm-thick gel cushion for sonography, in order to have a basic absorption comparable to the in-vivo-situation, and X-rayed (focus-film distance: 1m, exposure 52/2.5). The X-ray-visible elements in the form of the clips were easily visible in the X-ray image. They could not separated from the mesh by gentle manual pulling.

Example 3

Manufacture of an X-ray-visible, fine and coarse-pored polypropylene mesh with clips made from implant steel

The procedure was as in Example 2, but instead of the small clips, implantable clips of medium size made from stainless steel were used ("LIGACLIP Extra", medium-size, made from stainless steel with the US-material number 316L, from Ethicon Endo-Surgery). The clips could be very well recognised in the X-ray image.

Example 4

Manufacture of a mesh which is X-ray-visible, fine and coarse-pored, partly-resorbable and reinforced with medical steel tubes

Sleeves made from a medical chromium/nickel steel tube each with a length of 4 mm, an external diameter of 1.3 mm and an internal diameter of 1.0 mm were each fixed at 25 mm intervals, by compression at one end, on a monofilament (called "Pronova") manufactured by Ethicon Inc. which was 5.0 mil (0.127 mm) thick and made from fluorinated polyolefins. The fixed objects could not be moved on the monofilament manually. The monofilament prepared in this way was incorporated as a thread into an implant mesh customary in the trade ("Vypro", Ethicon GmbH) serving as a flexible basic structure and showed a very good contrast in the X-ray image.

Example 5

Manufacture of a polypropylene mesh which is X-ray-visible, fine and coarse-pored, reinforced with steel tubes made from non-magnetic implant steel

5

Steel sleeves made from non-magnetic implant steel ("Phynox", from Minitubes, Grenoble) each with a length of 4 mm, an external diameter of 1.2 mm and an internal diameter of 0.8 mm were fixed, by knotting in front of and behind each sleeve, on a thread made from polypropylene ("Prolene", 6.0 mil (0.1524 mm) diameter, Ethicon Inc.). The distance between each of the sleeves was approx 30 mm. These threads were drawn in between every fifth and sixth wale into a polypropylene mesh customary in the trade and showed a very good contrast in the X-ray image behind an approx. 5 cm-thick gel cushion.

15

Example 6

Marking of a polypropylene mesh with welded pre-shaped bodies made from zirconium dioxide as X-ray-visible elements

20

Balls made from stabilized zirconium dioxide (diameter 1.5 mm, YSZ-Ytt-stabilized, 95% ZrO_2 , 5% Y_2O_3 , Mühlmeier Mahltechnik) were placed on a polypropylene mesh customary in the trade ("Prolene-Netz", Ethicon GmbH) serving as basic structure, each at 3 cm intervals. Subsequently, circular pieces with a diameter of approx 1 cm were cut out of a 250 μ m-thick polypropylene foil, and were pressed centered onto the balls and each welded with 8 to 10 weld points with a "USG 440" ultrasound welding probe manufactured by Lühr. Subsequently, foil pieces of the same size were welded onto the back of the mesh as counterpieces.

25

30

The pre-shaped bodies made from stabilized zirconium dioxide were very clearly visible in the X-ray image (very good contrast behind a 10 cm-thick gel cushion) and could not be separated

35

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from the implant mesh by manual pulling on the implant mesh, rubbing or repeated bending.

Example 7

- 5 Marking of a polypropylene mesh with welded pre-shaped bodies made from zirconium dioxide as X-ray-visible elements

The procedure was as in Example 6, but in contrast to this, balls with a diameter of 0.5 mm were used (YSZ-Ytt-stabilized,
10 95% ZrO_2 , 5% Y_2O_3 , Mühlmeier Mahltechnik), and a 50 μm -thick polypropylene foil was used as a foil. A still very good contrast was shown in the X-ray image, but slightly fainter than in Example 6.

15 Example 8

Manufacture of a polypropylene mesh with X-ray-visible symbols

Symbols in the shape of a lower-case letter "e" (each 5 mm long, 5 mm wide and approx. 0.5 mm thick) were sewn onto a mesh customary in the trade, made from polypropylene as a flexible basic
20 structure ("Prolene", Ethicon GmbH), with a 0.127 mm-thick gold thread (Aldrich), each at 2.5 cm intervals. The X-ray-visible elements in the form of these letters could be well recognised in the x-ray image under a 10 cm-thick gel cushion.

25

Example 9

Marking of a polypropylene mesh with tube-reinforced balls made from zirconium dioxide

- 30 20 balls made from zirconium dioxide (0.5 mm diameter, YSZ-Ytt-stabilized, 95% ZrO_2 , 5% Y_2O_3 , Mühlmeier Mahltechnik) were attached by means of a polyethylene reinforcement to a polypropylene mesh customary in the trade ("Prolene", Ethicon GmbH) as a basic structure which was cut to a size of 6 cm x 7 cm. To this end,
35 each ball was packed centered into a 1.5 mm-long tube made from polyethylene with an external diameter of 1 mm and an internal

- 15 -

diameter of 0.5 mm from Portex (England). The tube pieces were arranged on the mesh so that each interval between the individual balls was 1.5 cm. Subsequently, the mesh was kept in a heatable printing press with baking paper for 10 to 30 seconds at a temperature of 135°C and a pressure of 2 bar.

Afterwards, the balls were encapsulated in rounded film pieces and firmly anchored to the mesh so that they could not be removed by rubbing or tearing until the mesh was destroyed. Figure 10 2 illustrates a section of the prepared implant. The mesh (basic structure) is designated by 20 and the zirconium dioxide balls by 22. The resulting rounded film pieces are indicated by the dots 24 in the area surrounding the balls 22.

15 In the case of the mesh treated under the conditions described, no thermal shrinkage was noticed. The mesh had the same flexural strength and tensile strength as the original mesh not subjected to an increased temperature. The increase in weight caused by the zirconium dioxide balls serving as X-ray-visible elements 20 was only 7%. The balls were very well visible in the X-ray image.

Example 10

Marking of an implant with encapsulated barium sulphate as an X-ray-visible substance

A 3.3 cm-long polypropylene tube (Portex) with an internal diameter of approx. 1.5 mm was filled with an approx 1 mm-thick barium sulphate-containing multifilament thread, (taken from the 30 swab material "Telacomp" from Hartmann) with a length of 2.7 cm so that there was a tube overhang of approx 3 mm to the left and to the right. This tube was bent twice so that the shape of an angular letter "C" resulted, and was fixed with bees wax on a commercially available polypropylene mesh ("Prolene-Netz", 6 mil 35 (0.1524 mm), Ethicon GmbH) and subsequently thermally welded, as described in Example 9.

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An almost round membrane with a diameter of approx 2 cm developed, in the centre of which the "C" made from the multifilament filled with barium sulphate could be recognised. The coating remained stable even when forcefully and rapidly bent and rubbed with the fingernails. The X-ray-visible element in the form of the "C" showed a good X-ray contrast.

Example 11

Marking of a partly-resorbable mesh with zirconium dioxide and subsequent hydrolysis

10 balls made from stabilized zirconium dioxide (0.5 mm diameter, YSZ-Ytt-stabilized, 95% ZrO_2 , 5% Y_2O_3 , Mühlmeier Mahltechnik) were attached via a polyethylene reinforcement to a partly-resorbable polypropylene-containing mesh ("Vypro-Netz", 15 cm x 10 cm, Ethicon GmbH) as a flexible basic structure. To this end, each ball was centred in an approx. 2 to 3 cm-long polyethylene tube (Portex 800/110/160) with an external diameter of 1 mm and an internal diameter of approx. 0.5 mm. The tube pieces were arranged on the mesh so that each interval between the balls was 2.5 cm; each tube piece was placed diagonally over 6 wales and 24 stitch rows. Subsequently, the mesh was kept in a heatable printing press with baking paper for 30 seconds at a temperature of 137 °C and a pressure of 2.2 bar.

The balls were then encapsulated in rounded film pieces and firmly anchored to the mesh so that they could not be removed by gentle rubbing or tearing. Furthermore, the mesh showed no thermal shrinkage under these conditions. The mesh had almost the same flexural strength and tensile strength as the original mesh not subjected to a thermal treatment. The zirconium dioxide markings were very clearly visible in the X-ray image. They could not be removed from the mesh by gentle rubbing and repeated bending.

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Subsequently, the resorbable part of the mesh was removed by being boiled for half an hour in a 10% soda solution. The resulting polypropylene mesh was rinsed with water and air-dried. The zirconium dioxide balls were still on the mesh and could not be removed from the mesh by gentle manual rubbing and repeated bending.

Example 12

Marking of a partly-resorbable mesh with zirconium dioxide in polypropylene reinforcement

The procedure was analogous to Example 11 but with the difference that a polypropylene tube (Portex "PP 60 x 100 FT", Lot G0515) served to reinforce the zirconium dioxide balls and the heatable printing press was kept for 3 minutes at a temperature of 121 °C and a pressure of 3 bar. The mesh showed practically no thermal shrinkage under these conditions. It had the same flexural strength and tensile strength as the original mesh. The markings in the form of zirconium dioxide balls could be very clearly seen in the X-ray image.

Example 13

Marking of a coarse-pored mesh with fine-pored marked mesh pieces

Circular pieces with a diameter of 1.4 cm were cut out round the zirconium dioxide balls from the mesh manufactured in Example 9. These round and small-pored mesh pieces were subsequently placed on a large-pored polypropylene mesh hydrolysed according to Example 1 (original material "Vypro", see Example 1) and welded with a "USG 440" type Lühr ultrasound welding probe with 8 to 10 weld points in each case.

Example 14

Manufacture of a polymer mesh which contains X-ray-visible polymer threads

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A polypropylene mesh was manufactured on a "RD3MT3/420 SN" type Müller crochet galloon machine as illustrated in Figure 3. Figure 3 also shows the corresponding pattern template. Like a conventional "Vypro" mesh manufactured by Ethicon GmbH, this mesh
5 was prepared with two part-wefts 32, 33, however the open-pillar stitches technique was chosen for the warp numbered 31.

In addition, several X-ray-visible stationary threads 34 were incorporated into the mesh, which were prepared as multifilament
10 polypropylene threads (300 tex) with a zirconium dioxide content of 50 wt.-%. The stationary threads 34 ran in the rapport of 10 wales between the two part-wefts 32 and 33 of the basic structure. As a meshing of these threads with the stationary threads 34 did not take place, the X-ray-visible stationary threads 34 were
15 not responsible for the stretching and strength properties of the implant.

Example 15

Manufacture of an X-ray-visible knitted product

20

A right/right-knitted product was manufactured on a Shima Seiki 12-pitch flatbed-knitting machine basically from a 320-den polypropylene yarn. An X-ray-visible thread was incorporated into every 11th stitch row. Each X-ray-visible thread, which was
25 prepared from a polypropylene yarn (300 tex) containing 35 wt.-% zirconium dioxide pigment, therefore follows on 10 stitch rows 40 (see Figure 4) in an incorporated loop 42.

By using the different threads and the chosen pattern, the knitted fabric acquired a transverse texture, however with sufficient elasticity.
30

Example 16

Marking of a partly-resorbable mesh with zirconium dioxide balls
35 in cords

- 19 -

A cord was braided on a customary ball-braiding machine with 36 bobbins from a polypropylene yarn (2 x 60 den polypropylene multifilament yarn per bobbin) with approx. 60 braids/inch. Zirconium dioxide balls each with a diameter of 1.5 mm were
5 incorporated therein at a mutual interval of approx. 10 mm (zirconium dioxide balls as in Example 6). After thermofixing of the braided product, the cord was incorporated as stationary thread into a composite mesh consisting of "Vicryl"- and polypropylene fibres. ("Vicryl" is a trade name of Ethicon GmbH for a resorba-
10 ble copolymer made from glycolide and lactide in a ratio of 9 to 1).

The implant obtained showed an only slightly higher flexural strength than an unaltered "Vypro" mesh. The zirconium dioxide
15 balls of the implant were very clearly visible in the X-ray image.

Example 17

Marking of a non-resorbable mesh with cords made from threads
20 filled with zirconium dioxide

A thread was manufactured on a customary ball braiding head with 8 bobbins from polypropylene multifilament yarn (60 den polypropylene multifilament yarn per bobbin) with approx. 60 braids/-
25 inch, into which zirconium dioxide balls with a diameter of 0.5 mm (zirconium dioxide balls as in Example 7) were incorporated, each at a 5 mm interval. After thermofixing the braided thread, the braided threads were incorporated into a knitwear product made from a non-resorbable, synthetic monofilament with a diame-
30 ter of approx. 0.1 mm.

The implant showed no noticeable change in flexural strength compared with a knitted mesh without an incorporated X-ray-visible thread. The zirconium dioxide balls were clearly visible in
35 the X-ray image.

- 20 -

Example 18

Marking of a non-resorbable mesh with knotted threads filled with zirconium dioxide

- 5 A thread, manufactured as in Example 17, however with only approx. 35 braids/inch, was knotted both in front of and behind each zirconium dioxide ball, in order on the one hand to prevent a displacement of the small balls, but on the other hand also to obtain a very soft braided thread. Subsequently, this very soft
10 braided thread was incorporated into a knitted fabric as in Example 17 without previous thermofixing.

The flexural strength and elasticity of the implant was not measurably altered by the thread filled with zirconium dioxide
15 balls. The zirconium dioxide balls placed in the implant were clearly visible in the X-ray image.

Example 19

Manufacture of a tape marked with X-ray-visible elements

20

- A so-called striped braid was manufactured on a customary braiding head, threads filled with approx. 30 wt.-% zirconium dioxide being braided round as so-called cores. For the braid construction, the braid had 19 bobbins each with 3 x 60 den polypropylene multifilament yarn. The six cores each consisted of a
25 polypropylene monofilament with a diameter of approx. 0.2 mm filled with approx. 30 wt.-% zirconium dioxide.

- The tape obtained was approx. 6 mm wide, and the X-ray image
30 showed the cores filled with the X-ray-visible zirconium dioxide as six fine bands.

Claims

1. Areal implant with a flexible polymer-based basic structure
(10; 20; 31, 32, 33; 40) and with X-ray-visible elements
5 (12; 22; 34; 42).
2. Implant according to claim 1, characterized in that the X-
ray-visible elements (12; 22; 34; 42) are arranged in an
areal pattern.
10
3. Implant according to claim 1 or 2, characterized in that the
basic structure (10; 20; 31, 32, 33; 40) includes non-resor-
bable polymer.
- 15 4. Implant according to one of claims 1 to 3, characterized in
that the basic structure (10; 20; 31, 32, 33; 40) has one of
the forms chosen from the following group: meshes, tapes,
foils, perforated foils.
- 20 5. Implant according to one of claims 1 to 4, characterized in
that the X-ray-visible elements (12; 22; 34; 42) have at
least one of the X-ray-visible substances chosen from the
following group: pure zirconium dioxide, stabilized zirco-
nium dioxide, zirconium nitride, zirconium carbide, tanta-
25 lum, tantalum pentoxide, barium sulphate, silver, silver io-
dide, gold, platinum, palladium, iridium, copper, ferric
oxides, not very magnetic implant steels, non-magnetic im-
plant steels, titanium, alkali iodides, iodated aromatics,
iodated aliphatics, iodated oligomers, iodated polymers,
30 alloys of substances thereof capable of being alloyed.
6. Implant according to one of claims 1 to 5, characterized in
that at least part of the X-ray-visible elements is formed
as pre-shaped bodies (12; 22) of respective length, width
35 and height in the range of 0.1 mm to 50 mm, the pre-shaped

- 22 -

bodies (12; 22) being attached to the basic structure (10; 20).

- 5 7. Implant according to claim 6, characterized in that at least one pre-shaped body comprises a mixture of at least one X-ray-visible substance with a binding agent, preferably a non- or slowly resorbable polymer and/or a wax.
- 10 8. Implant according to claim 6 or 7, characterized in that at least one pre-shaped body (12; 22) has one of the shapes chosen from the following group: beads, balls, small tubes, rods, small plates, rings, discs, bones, clips.
- 15 9. Implant according to one of claims 6 to 8, characterized in that at least one pre-shaped body (22) has a coating and/or outer layer (24) made from non-resorbable material.
- 20 10. Implant according to one of claims 6 to 9, characterized in that at least one pre-shaped body (12; 22) is attached to the basic structure or a holding device connected to the basic structure by knotting, compression, welding and/or gluing.
- 25 11. Implant according to one of claims 1 to 10, characterized in that at least part of the X-ray-visible elements comprises a polymer tube or a cord which is filled at least partly with particles of a size of at most 2.5 mm made from an X-ray-visible substance.
- 30 12. Implant according to claim 11, characterized in that the X-ray-visible substance is additionally fixed in the polymer tube or the cord, preferably by thermal shrinking and/or gluing.
- 35 13. Implant according to one of claims 1 to 12, characterized in that at least part of the X-ray-visible elements (34; 42) is

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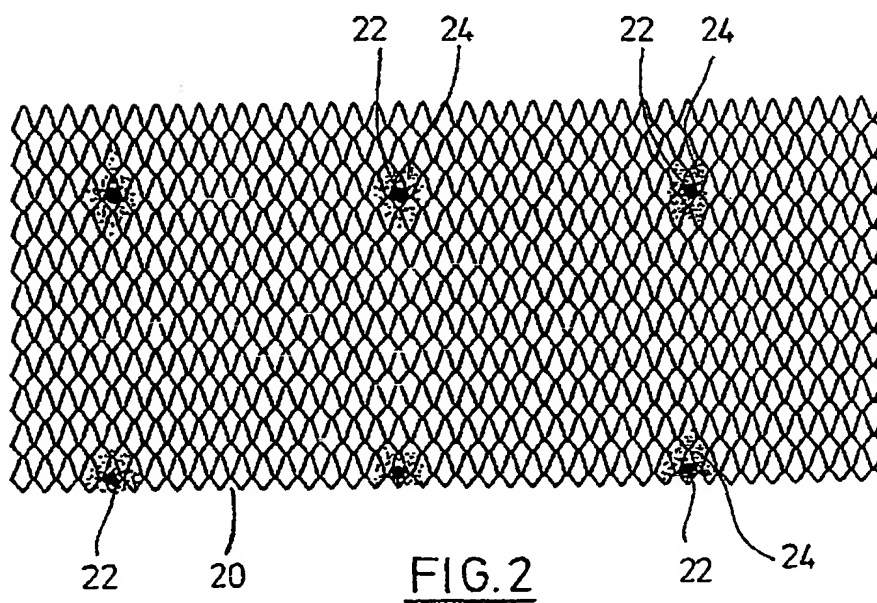
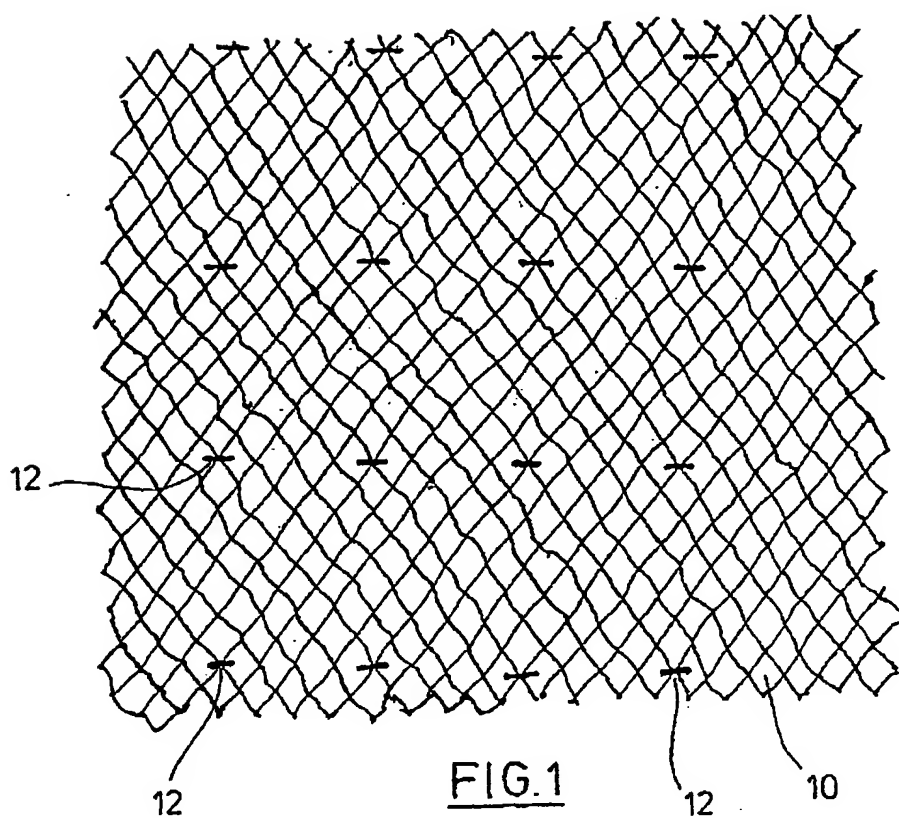
designed as polymer, into which an X-ray-visible substance made from particles of a size in the range of 10 nm to 500 μm , preferably 10 nm to 100 μm , is compounded.

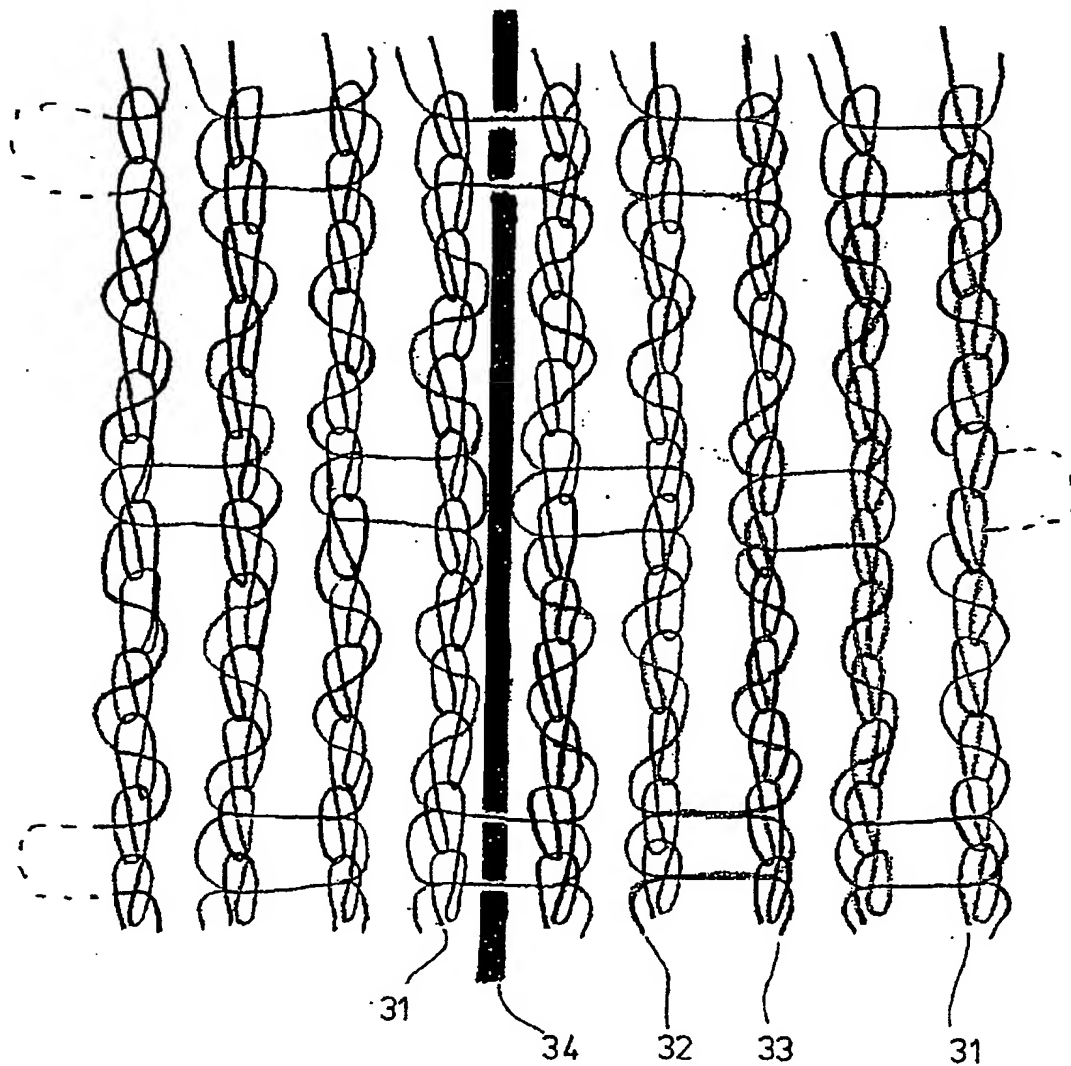
- 5 14. Implant according to claim 13, characterized in that the polymer with X-ray-visible substance (34; 42) is present in the implant in at least one of the forms chosen from the following group: monofilaments, monofilaments reduced along sections, multifilaments, twines, braided threads, cords,
10 foils, films, small film tapes, tapes, such forms which are provided with knots.
- 15 15. Implant according to claim 14, characterized in that the basic structure (40) has polymer with X-ray-visible substance (42).
- 20 16. Implant according to claim 14 or 15, characterized in that polymer with X-ray-visible substance (34) is attached as additional component to the basic structure (31, 32, 33).
17. Implant according to one of claims 13 to 16, characterized in that the X-ray-visible substance of the polymer comprises zirconium dioxide.
- 25 18. Implant according to one of claims 13 to 17, characterized in that the X-ray-visible substance of the polymer comprises barium sulphate and this polymer is additionally coated with a non-resorbable polymer or wax.
- 30 19. Implant according to one of claims 1 to 18, characterized in that at least part of the X-ray-visible elements comprises an X-ray-visible metal thread.
- 35 20. Implant according to one of claims 1 to 19, characterized in that at least part of the X-ray-visible elements comprises

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an X-ray-visible symbol, which is preferably provided repeatedly and at equal intervals.

21. Implant according to claim 20, characterized in that the
5 symbol is designed in one of the forms chosen from the following group: sewn from X-ray-visible threads, stitched from X-ray-visible threads, embossed from X-ray-visible foil, put together from X-ray-visible objects, put together from X-ray-visible powder.
- 10 22. Implant according to one of claims 1 to 21, characterized in that the implant contains at least one of the substances chosen from the following group: polyalkenes, polypropylene, polyethylene, fluorinated polyolefins, polytetrafluoroethylene,
15 polyvinylidene fluoride, polyamides, polyurethanes, polyisoprenes, polystyrenes, polysilicones, polycarbonates, polyaryletherketones, polymethacrylates, polyacrylates, aromatic polyesters, polyimides, copolymers of polymerisable substances thereof.
- 20 23. Implant according to one of claims 1 to 22, characterized in that the basic structure has a proportion of resorbable polymer, which preferably contains at least one of the substances chosen from the following group: polyhydroxy acids,
25 polylactides, polyglycolides, polyhydroxybutyrates, polyhydroxyvalerates, polycaprolactones, polydioxanones, synthetic and natural oligo- and polyaminoacids, polyphosphazenes, polyanhydrides, polyorthoesters, polyphosphates, polyphosphonates, polyalcohols, polysaccharides, polyethers,
30 resorbable glasses, copolymers of polymerisable substances thereof.
- 35 24. Implant according to one of claims 1 to 23, characterized in that the implant is also detectable by means of ultrasound and/or magnetic resonance tomography.

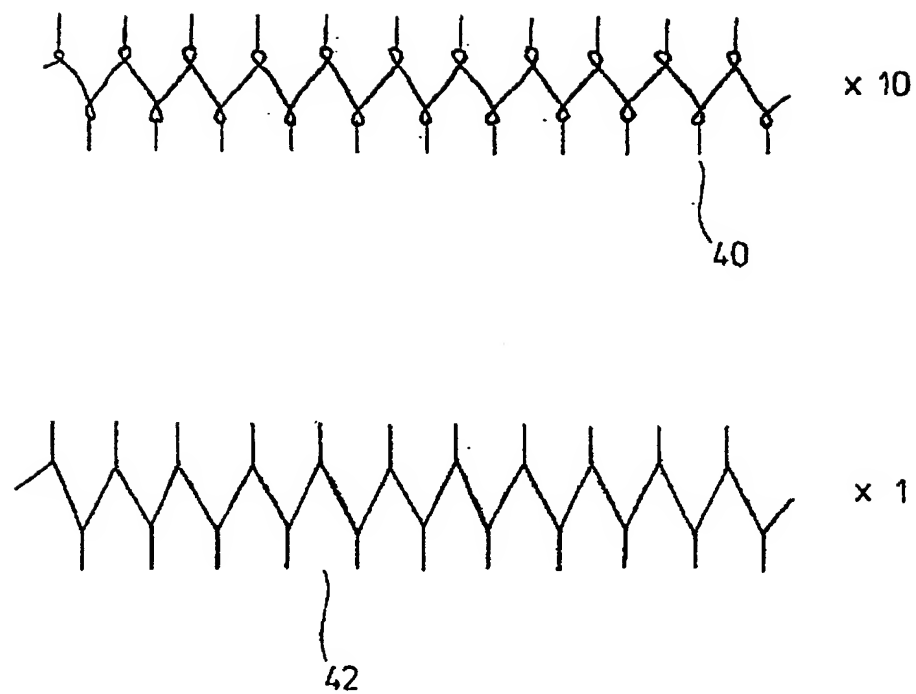




Pattern Template – Example 14

warp (31)	part-weft I (32)	thread (34)	part-weft II (33)
0 2 2 0	0 4 2 4 2 6 2 4 2 4	0 0	6 2 4 2 4 0 4 2 4 2

FIG.3

FIG. 4

INTERNATIONAL SEARCH REPORT

International application No

PCT/EP 01/00122

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61F2/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	FR 2 712 177 A (SGRO) 19 May 1995 (1995-05-19)	1,2,4,5, 19,22,24
Y	page 2, line 11 - line 13; figures	3,6,8, 10, 13-15, 20,21,23
Y	EP 0 797 962 A (ETHICON GMBH & CO. KG) 1 October 1997 (1997-10-01) the whole document	3,23
Y	EP 0 783 873 A (BOSTON SCIENTIFIC TECHNOLOGY INC.) 16 July 1997 (1997-07-16) column 12, line 45 - line 56; figure 4A	6,8,10, 20,21
Y	WO 90 03036 A (SMID ET AL) 22 March 1990 (1990-03-22) the whole document	13-15

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

In relation on patent family members

International application No

PCT/EP 01/00122

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
FR 2712177 A	19-05-1995	NONE	
EP 797962 A	01-10-1997	DE 19613730 A	02-10-1997
		AU 704334 B	22-04-1999
		AU 1648597 A	02-10-1997
		BR 9701466 A	10-11-1998
		CA 2200801 A	26-09-1997
		JP 10024054 A	27-01-1998
		US 6162962 A	19-12-2000
		ZA 9702563 A	25-09-1998
EP 783873 A	16-07-1997	US 5609627 A	11-03-1997
		GR 3033808 T	31-10-2000
		DE 29521548 U	10-07-1997
		DE 29521776 U	27-08-1998
		DE 29522113 U	07-10-1999
		DE 29522160 U	13-04-2000
		DE 29522161 U	30-03-2000
		DE 29522162 U	30-03-2000
		EP 0782841 A	09-07-1997
		EP 0783874 A	16-07-1997
		GR 3033211 T	31-08-2000
		AT 188604 T	15-01-2000
		AT 191628 T	15-04-2000
		AT 191629 T	15-04-2000
		AU 693527 B	02-07-1998
		AU 1870995 A	29-08-1995
		AU 3224599 A	05-08-1999
		AU 719206 B	04-05-2000
		AU 6381598 A	18-06-1998
		AU 717678 B	30-03-2000
		AU 6381698 A	18-06-1998
		CA 2182982 A	17-08-1995
		DE 69514511 D	17-02-2000
		DE 69514511 T	05-10-2000
		DE 69516292 D	18-05-2000
		DE 69516292 T	14-12-2000
		DE 69516293 D	18-05-2000
		DE 69516293 T	31-08-2000
		DK 759729 T	08-05-2000
		DK 783873 T	18-09-2000
		DK 783874 T	24-07-2000
		EP 0759729 A	05-03-1997
		ES 2141339 T	16-03-2000
		ES 2146951 T	16-08-2000
		ES 2144824 T	16-06-2000
		GR 3033133 T	31-08-2000
		JP 9511160 T	11-11-1997
		US 6117167 A	12-09-2000
		WO 9521592 A	17-08-1995
		US 6165213 A	26-12-2000
		US 5716365 A	10-02-1998
		US 5693086 A	02-12-1997
		US 5776180 A	07-07-1998
		US 5800508 A	01-09-1998
		US 5718724 A	17-02-1998
		US 5683450 A	04-11-1997
		US 5916263 A	29-06-1999